AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

- 1. (currently amended) A method for predicting an increased risk for onset of glaucoma in a subject, comprising assaying for the presence of A gene assay method comprising the steps of: detecting a mutation of at least one nucleotide base in the coding region of an optineurin (OPTN) gene of said a human subject, wherein when said mutation is present, said subject is predicted to have an increased risk for onset of glaucoma; and predicting future onset of glaucoma in the subject using the mutation as an index.
- 2. (currently amended) The gene assay method of claim 1, wherein the coding region of said glaucoma related gene is an OPTN gene comprises nucleotides 1 to 1734 of has a nucleic acid sequence denoted by SEQ ID NO: 1.
- 3. (currently amended) The gene assay method of claim 2, wherein said mutation is a substitution of G for A at position 619 and/or a substitution of A for G at position 898, or both, in the nucleic acid sequence denoted by SEQ ID NO:1.
- 4. (currently amended) The gene assay method of claim 2, wherein said mutation is a deletion of one or more <u>nucleotides in bases in the nucleic acid sequence denoted by SEQ ID</u>
 NO: 1.
- 5. (currently amended) The gene assay method of claim 2, wherein said mutation is an insertion of one or more <u>nucleotides in bases in the nucleic acid sequence denoted by SEQ ID</u>
 NO: 1.

- 6. (currently amended) The gene assay method of claim 2, wherein said mutation is two or more substitutions of <u>nucleotides in bases in the nucleic acid sequence denoted by SEQ</u>

 ID NO: 1.
- 7. (currently amended) The gene assay method according to claim 1, wherein the said glaucoma is primary open angle glaucoma and/or normal ocular tension glaucoma, or both.
- 8. (currently amended) The gene assay method according to claim 1, wherein the said assay is carried out mutation is detected by using an oligonucleotide that hybridizes to a selected portion capable of forming a hybrid at a specific position of the coding region of the OPTN gene.
 - 9. (canceled).
- 10. (currently amended) A gene assay method for predicting an increased risk for future onset of primary open angle glaucoma and/or normal ocular tension glaucoma, or both, in a subject, comprising the steps of:
- (a) isolating a polynucleotide sample from a subject suspected of having a mutation in a glaucoma related gene,
- (ab) performing a nucleic acid amplification process on a polynucleotide sample from a subject said polynucleotide using at least one oligonucleotide primer pair, wherein said primer oligonucleotide pair comprises a member selected from the group consisting of oligonucleotides comprising sequences as follows:
 - (1) an oligonucleotide <u>primer pair</u> consisting of <u>nucleotide sequences a base</u> sequence represented by (i) 21 and 22, and (ii) 27 and 28 any of SEQ ID NOs: 15 to 40;

- (2) <u>an oligonucleotide primer pair wherein each member of said pair is a</u>

 <u>complement of one member a complementary chain of an oligonucleotide primer pair</u>

 according to (1);
- (3) an oligonucleotide <u>primer pair wherein each member of said pair that</u>
 hybridizes with <u>one member of an oligonucleotide primer pair according to (1) or (2) under stringent conditions;</u>
- (4) an oligonucleotide <u>primer pair wherein each member of said pair has</u>

 having a homology of 60% or more to a respective member of to an oligonucleotide

 <u>primer pair according to any one of (1) to (3)-; and</u>
- (5) an oligonucleotide <u>primer pair</u> according to any one of (1) to (4) <u>wherein</u> each member of said pair has at least one nucleotide <u>having one to several bases mutated</u> by substitution, deletion, insertion or addition <u>mutation</u>,
- (be) detecting a mutation of at least one base nucleotide in the coding region of the amplification product of (a)a glaucoma related gene; and
- increased risk for onset of primary open angle glaucoma or normal ocular tension glaucoma, or both predicting future onset of primary open angle glaucoma and/or normal ocular tension glaucoma glaucoma using the mutation as an index.
 - 11. (canceled).
- 12. (new) The method according to claim 10, wherein said at least one oligonucleotide primer pair amplifies a selected portion of the coding region of an OPTN gene comprising nucleotides 1 to 1734 of SEQ ID NO: 1

13. (new) The method of claim 12, wherein said mutation is a substitution of G for A at position 619 or a substitution of A for G at position 898, or both, in SEQ ID NO:1.